Serial No. 08/484,542



Remarks

Claims 56 and 57 have been amended in this supplemental amendment. Claims 27-35 and 56-60 are active in this application.

Support for the amendment of claim 56 is found in the specification at page 8, line 19. Support for the amendment of claim 57 is found in the specification at page 10, lines 5-7. No new matter has been added.

Provided herewith is a version of amended claims 56 and 57, with markings to show the changes made.

In the amendment and reply filed October 9, 2001, claims 29, 31 and 33 were amended. In the version of the amended claims with markings to show changes made that was filed with the amendment, amended claim 29 was inadvertently numbered incorrectly as claim 30, and amended claim 33 was inadvertently numbered incorrectly as claim 34. Accordingly, provided herewith is a version of correctly numbered amended claims 29, 31 and 33, with markings to show changes made.

Respectfully submitted,

Attorney for Applicant Registration No. 41,264 317-276-1664

Eli Lilly and Company Patent Division/DC 1501 Lilly Corporate Center Indianapolis, Indiana 46285

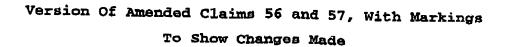
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- 56. (Once Amended) The composition of Claim 35, wherein the phenolic compound [preservative] is selected from the group consisting of phenol and m-cresol.
- 57. (Once amended) The composition of claim 28, wherein the composition is a pharmaceutical composition that further comprises a phenolic compound, glycerol, and a pharmaceutically acceptable buffer.

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Serial No. 08/484,542

V rsion Of Amended Claims 29, 31 and 33 (Amended October 9, 2001), With Markings To Show Changes Made

- 29. (Once Amended) The composition of Claim 28, wherein the solution comprises [further comprising from] about 0.2 mole to about 0.7 mole of zinc per mole of fatty acidacylated insulin.
- 31. (Once Amended) The composition of Claim 30, further comprising a phenolic compound at a concentration of from 0.5 mg to 5 mg [of] per milliliter of the aqueous solution.
- 33. (Once Amended) The composition of claim 32, wherein the fatty acid-acylated insulin is N-palmitoyl LysB29 human insulin, and wherein the solution comprises [is comprised of] from about 0.3 mole to about 0.55 mg per milliliter of the aqueous solution.

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Legal Department - Patent Division

Marianne P. Allen, Primary Examiner

Company: USPTO Fax: 1-703-305-3014 Phone: 1-703-308-0666 Date: December 11, 2001

From: Grant E. Reed Fax: 317-276-5172 Phone: 317-276-1664

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Re:

Applicant: Mark Brader, et al.

Serial No.: 08/484,542 Filed: June 7, 1995

For: Stabilized, Acylated Insulin Formulations

Our Docket No.: X-10097

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